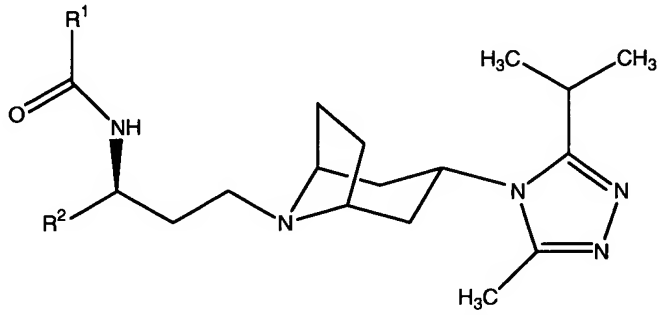
Listing of Claims

Claims 1 - 37 (canceled).

Claim 38 (currently amended): A method of ~~treating~~ antagonizing a CCR5 receptor in a mammal ~~a disorder in which the modulation of CCR5 receptors is implicated~~ comprising administering to said mammal an effective amount of a compound of Formula I



I

wherein R^1 is C_{3-6} cycloalkyl optionally substituted by one or more fluorine atoms, or C_{1-6} alkyl optionally substituted by one or more fluorine atoms, or C_{3-6} cycloalkylmethyl optionally ring-substituted by one or more fluorine atoms, and

R^2 is phenyl optionally substituted by one or more fluorine atoms,

or a pharmaceutically acceptable salt or solvate thereof.

Claim 39 (previously presented): A method of treating in a mammal a respiratory disorder selected from adult respiratory distress syndrome (ARDS), bronchitis, chronic bronchitis, chronic obstructive pulmonary disease, cystic fibrosis, asthma, emphysema, rhinitis and chronic sinusitis, which comprises administering to said mammal an effective amount of a compound of Formula I according to claim 38.

Claim 40 (currently amended): A method of treating in a mammal an inflammatory bowel disease, multiple sclerosis, rheumatoid arthritis, graft rejection, ~~including a kidney or a lung allograft,~~ endometriosis, type I diabetes, a renal disease, chronis pancreatitis, an inflammatory lung condition or chronic heart failure which comprises administering to said mammal an effective amount of a compound of Formula I according to claim 38.

Claim 41 (previously presented): A method of treating HIV infection comprising administering an effective amount of a compound of Formula I according to claim 38.

Claim 42 (previously presented): A method of treating HIV infection comprising administering an effective amount of a compound selected from the group consisting of:

N-{(1S)-3-[3-(3-Isopropyl-5-methyl-4H-1,2,4-triazol-4-yl)-exo-8-azabicyclo[3.2.1]oct-8-yl]-1-phenylpropyl}cyclobutanecarboxamide;

N-{(1S)-3-[3-(3-Isopropyl-5-methyl-4H-1,2,4-triazol-4-yl)-exo-8-azabicyclo[3.2.1]oct-8-yl]-1-phenylpropyl}cyclopentanecarboxamide;

N-{(1S)-3-[3-(3-Isopropyl-5-methyl-4H-1,2,4-triazol-4-yl)-exo-8-azabicyclo[3.2.1]oct-8-yl]-1-phenylpropyl}-4,4,4-trifluorobutanamide;

N-{(1S)-3-[3-(3-Isopropyl-5-methyl-4H-1,2,4-triazol-4-yl)-exo-8-azabicyclo[3.2.1]oct-8-yl]-1-phenylpropyl}-4,4-difluorocyclohexanecarboxamide; and

N-{(1S)-3-[3-(3-Isopropyl-5-methyl-4H-1,2,4-triazol-4-yl)-exo-8-azabicyclo[3.2.1]oct-8-yl]-1-(3-fluorophenyl)propyl}-4,4-difluorocyclohexanecarboxamide;

or a pharmaceutically acceptable salt or solvate thereof.

Claim 43 (previously presented): The method of claim 40 wherein said graft rejection is kidney or lung allograft rejection.

Claim 44 (currently amended): A method according to claim 38 wherein said ~~modulation~~ antagonizing comprises reducing or inhibiting the CCR5 receptor-associated responses in said mammal.

Claim 45 (new): A pharmaceutical composition comprising a compound of Formula I according to claim 38 or a pharmaceutically acceptable salt or solvate thereof together with one or more pharmaceutically acceptable excipients, diluents or carriers, and one or more additional therapeutic agents.